

UNITED STATES DISTRICT COURT
SOUTHERN DISTRICT OF CALIFORNIA

KATHLEEN HOLT and JOSE
RUVALCABA,

Plaintiff,

v.

FOODSTATE, INC., dba MEGAFOOD,
dba INNATE RESPONSE FORMULAS,

Defendant.

Case No.: 15cv78 L (JMA)

**ORDER GRANTING IN PART AND
DENYING IN PART DEFENDANT'S
MOTION TO DISMISS [ECF No. 21]
and GRANTING LEAVE TO AMEND**

Currently pending is defendant's motion to dismiss plaintiffs' first amended complaint. The motion has been fully briefed and is considered without oral argument.

I. Background

Defendant, Foodstate, Inc. ("Foodstate"), is a producer in the United States of health supplements including the multivitamins at issue. In this case, Plaintiffs Kathleen Holt and Jose Ruvalcaba ("Plaintiffs") purchased and consumed different vitamin products that Defendant produces, One Daily Multivitamin and Men's One Daily. (First Amended Complaint "FAC" ¶¶ 29, 33, 35, 39.) Plaintiffs allege, on behalf of themselves and all others similarly situated, that Defendant falsely represents that its products are entirely or largely from whole food sources, when in fact they are also derived from

1 synthetic sources. Next, Plaintiffs argue that the products contain “magnesium stearate,
2 calcium stearate, or any other stearate/stearic acid” that “may be harmful and undesirable
3 to consumers” and which are not on Defendant’s product’s labels. (FAC ¶ 42.)

4 Plaintiffs allege five causes of action in the FAC: (1): violation of California’s
5 False Advertising Law (“FAL”); (2) violation of California’s Sherman Law (“Sherman”);
6 (3) violation of California’s Unfair Competition Law (“UCL”); (4) negligent
7 misrepresentation; and (5) intentional misrepresentation. Foodstate now moves to dismiss
8 all claims based on lack of standing to bring claims for products Plaintiffs did not
9 purchase because the unpurchased products are not substantially similar to the products
10 they purchased. Foodstate also moves to dismiss claims based on express and implied
11 preemption and Plaintiffs’ alleged failure to meet Federal Rule of Civil Procedure 9(b)’s
12 heightened pleading standard for fraud-based claims.

13 **II. Legal Standard for Motion to Dismiss under Rule 12(b)(6)**

14 The court must dismiss a cause of action for failure to state a claim upon which
15 relief can be granted. FED. R. CIV. P. 12(b)(6). A motion to dismiss under Rule 12(b)(6)
16 tests the legal sufficiency of the complaint. *Navarro v. Block*, 250 F.3d 729, 732 (9th Cir.
17 2001). The court must accept all allegations of material fact as true and construe them in
18 light most favorable to the nonmoving party. *Cedars-Sinai Med. Ctr. v. Nat’l League of*
19 *Postmasters of U.S.*, 497 F.3d 972, 975 (9th Cir. 2007). Material allegations, even if
20 doubtful in fact, are assumed to be true. *Bell Atl. Corp. v. Twombly*, 550 U.S. 544, 555
21 (2007). However, the court need not “necessarily assume the truth of legal conclusions
22 merely because they are cast in the form of factual allegations.” *Warren v. Fox Family*
23 *Worldwide, Inc.*, 328 F.3d 1136, 1139 (9th Cir. 2003) (internal quotation marks omitted).
24 In fact, the court does not need to accept any legal conclusions as true. *Ashcroft v. Iqbal*,
25 129 S. Ct. 1937, 1949 (2009)

26 A complaint may be dismissed as a matter of law either for lack of a cognizable
27 legal theory or for insufficient facts under a cognizable theory. *Robertson v. Dean Witter*
28 *Reynolds, Inc.*, 749 F.2d 530, 534 (9th Cir. 1984). “While a complaint attacked by a Rule

1 12(b)(6) motion to dismiss does not need detailed factual allegations, a plaintiff's
2 obligation to provide the 'grounds' of his 'entitlement to relief' requires more than labels
3 and conclusions, and a formulaic recitation of the elements of a cause of action will not
4 do." *Twombly*, 550 U.S. at 555 (internal citations omitted). Instead, the allegations in the
5 complaint "must be enough to raise a right to relief above the speculative level." *Id.*
6 Thus, "[t]o survive a motion to dismiss, a complaint must contain sufficient factual
7 matter, accepted as true, to 'state a claim to relief that is plausible on its face.'" *Iqbal*, 129
8 S. Ct. at 1949 (citing *Twombly*, 550 U.S. at 570). "A claim has facial plausibility when
9 the plaintiff pleads factual content that allows the court to draw the reasonable inference
10 that the defendant is liable for the misconduct alleged." *Id.* "The plausibility standard is
11 not akin to a 'probability requirement,' but it asks for more than a sheer possibility that a
12 defendant has acted unlawfully." *Id.*

13 Generally, courts may not consider material outside the complaint when ruling on a
14 motion to dismiss. *Hal Roach Studios, Inc. v. Richard Feiner & Co.*, 896 F.2d 1542, 1555
15 n.19 (9th Cir. 1990). However, documents specifically identified in the complaint whose
16 authenticity is not questioned by parties may also be considered. *Fecht v. Price Co.*, 70
17 F.3d 1078, 1080 n.1 (9th Cir. 1995) (superceded by statutes on other grounds).
18 Moreover, the court may consider the full text of those documents, even when the
19 complaint quotes only selected portions. *Id.* It may also consider material properly
20 subject to judicial notice without converting the motion into one for summary judgment.
21 *Barron v. Reich*, 13 F.3d 1370, 1377 (9th Cir. 1994).

22 As a preliminary matter, Defendants request judicial notice of images of eight of
23 its products taken from Defendant's website. Plaintiff has not opposed the request.
24 Accordingly, the Court will grant Defendants' request and will consider the contents
25 therein. *See Bruton v. Gerber Prods. Co.*, 961 F.Supp.2d 1062, 1075, n. 1 (N.D. Cal.
26 2013) (granting request to take judicial notice of labels for ten products referenced in the
27 complaint).

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III. Discussion

A. Standing to Assert Claims as to Unpurchased Products

Plaintiff Holt purchased One Daily Multivitamin and Plaintiff Ruvalcaba purchased Men's One Daily. In their FAC, Plaintiffs also include 107 additional Foodstate products that they never purchased in their putative class definition. Foodstate moves to dismiss Plaintiffs' claims, arguing that Plaintiffs lack standing with respect to products they did not purchase. Plaintiffs counter that they have standing to bring claims for unpurchased products that are substantially similar to the products they purchased.

Standing under Article III and the UCL and FAL requires that a plaintiff suffer injury-in-fact. *Lujan v. Defenders of Wildlife*, 504 U.S. 555, 560 (1992). Plaintiffs may have standing to assert claims for unnamed class members based on products he or she did not purchase so long as the products *and* alleged misrepresentations are substantially similar. *Dorfman v. Nutramax Laboratories, Inc.*, 2013 WL 5353043, (S.D. Cal. Sept. 23, 2013) (emphasis added). Although Plaintiffs have not purchased 107 of the other vitamin/supplement products produced by Defendant, the allegations of misrepresentation mimic those made against One Daily Multivitamin and Men's One Daily. Specifically, Plaintiffs allege that 83 of the other products wrongfully purport to contain nutrients from whole food *and* the presence of magnesium stearate, calcium stearate or any other stearate/stearic acid was not disclosed. Additionally, Plaintiffs allege six of the other products wrongfully claim to contain nutrients from whole foods. Lastly, Plaintiffs allege 18 of the final 107 products do not disclose the presence of magnesium stearate, calcium stearate, or any other stearate/stearic acid. These three claims of misrepresentation overlap.

However, Plaintiffs have failed to show that the 107 products are substantially similar to the two purchased products in order to pass muster for standing. Although the ingredients are not required to be identical in order to be substantially similar, the products must contain similar ingredients and must have similar packaging and labeling. *Dysthe v. Basic Research LLC*, 2011 WL 5868307 *4-5 (C.D. Cal. 2011). In *Dysthe*, the

1 court found that two weight-loss products made by the same manufacturer were not
2 substantially similar because they contained significant different ingredients. *Dysthe*, WL
3 5868307 at *4. The first product contained 19 ingredients, while the second contained
4 ten. Although there was some overlap of ingredients, it did not amount to the standard
5 needed to satisfy standing. Additionally, both products contained different percentage
6 daily values of the ingredients within them. In the present case, there is overlap of some
7 ingredients in some of the products; however, Plaintiffs have failed to show that there is
8 substantial similarity amongst all of the 109 products.

9 The court in *Dysthe* also looked at the packaging to distinguish the products.
10 *Dysthe*, 2011 WL 5868307 at *5. The packaging of the products differed in color and
11 text. Similarly, the packaging in this case is significantly different between even the two
12 consumed products, let alone the 107 additional products. Because plaintiffs have not
13 alleged that the unpurchased products are substantially similar to the products plaintiffs
14 purchased, defendant's motion to dismiss based on lack of standing will be granted.
15 Plaintiffs will be given an opportunity, however, to amend their complaint to cure the
16 deficiencies noted if they are able to do so.

17 **B. Preemption**

18 Defendant argues that Plaintiffs' claims are expressly and impliedly preempted by
19 federal law. (Def.'s Mot. Dismiss 10.) Defendant first contends that the Nutrition,
20 Labeling, and Education Act ("NLEA"), a 1990 amendment to the Food, Drug and
21 Cosmetic Act ("FDCA"), expressly preempts Plaintiffs' claim. Further, Defendant argues
22 that Plaintiffs' state law fraud claims are barred by implied preemption because the
23 claims conflict with the FDCA's enforcement scheme and improperly seek to have the
24 Court play the role of the FDA.

25 **1. Express Preemption Under the FDCA**

26 Defendant contends that Plaintiffs' claims are expressly preempted by federal law.
27 (Def.'s Mot. Dismiss 10.) Defendant supports this contention by asserting the NLEA, an
28 amendment to the FDCA, expressly preempts Plaintiffs' attempt to impose food labeling

1 requirements not identical to those set forth under federal law. Defendant argues that
 2 Plaintiffs seek to impose liability on Defendant for omitting label information not required
 3 by federal law but instead, for Plaintiffs' "articulation of consumer preferences." (Def.'s
 4 Mot. Dismiss 11.) Specifically, Defendant contends that federal law allows magnesium
 5 stearate, calcium stearate or any other stearate/steric acid to be included in food as
 6 "incidental additives" if in "insignificant amounts." (*Id.* at 11.) 21 C.F.R. 101.100(a)(3)(ii)
 7 provides that "incidental additives that are present in a food at insignificant levels and do
 8 not have any technical or functional effect in that food" need not be included on the
 9 ingredient label. But Plaintiffs counter that their claims are consistent with requirements
 10 under federal law because the question is whether an additive is included at a significant
 11 level which is a factual question, better suited for summary judgment or trial. (Pls'. Opp'n.
 12 at 11.)

13 "It is well established that the party who asserts that a state law is preempted bears
 14 the burden of so demonstrating." *In re Farm Raised Salmon Cases*, 42 Cal. 4th 1077,
 15 1088 (2008). Such a presumption may be difficult to overcome when, "[i]n all pre-
 16 emption cases, and particularly in those in which Congress has 'legislated...in a field
 17 which the States have traditionally occupied,'" we "start with the assumption that the
 18 historic police powers of the States were not to be superseded by the Federal Act unless
 19 that was the clear and manifest purpose of Congress." *Medtronic, Inc. v. Lohr*, 518 U.S.
 20 470, 485 (1996) (quoting *Rice v. Santa Fe Elevator Corp.*, 331 U.S. 218, 230 (1947)).
 21 "[C]onsumer protection laws such as the [UCL], false advertising law, and CLRA, are
 22 within the states' historic police powers and therefore are subject to the presumption
 23 against preemption.' Laws regulating the proper marketing of food, including the
 24 prevention of deceptive sales practices, are likewise within states' historic police powers."
 25 *In re Farm Raised Salmon Cases*, 42 Cal. 4th at 1088 (quoting the appellate court, *In re*
 26 *Farm Raised Salmon Cases*, 48 Cal. Rptr. 3d 449, 453 (2006)).

27 Adding to this presumption against preemption, there is no indication that
 28 Congress intended the NLEA to preempt the state labeling requirements at issue in this

1 case. The congressional notes point to the opposite conclusion: “[t]he Nutrition Labeling
2 and Education Act [NLEA] of 1990 shall not be construed to preempt any provision of
3 State law, unless such provision is expressly preempted under section 403A of the
4 Federal Food, Drug, and Cosmetic Act [codified as 21 U.S.C.A. § 343–1].” NLEA,
5 PUB.L. NO. 101–535, § 6(c)(1), 104 Stat. 2353.

6 The NLEA section on preemption, 21 U.S.C.A. § 343–1, states that “except as
7 provided in subsection (b) of this section, no State... may directly or indirectly establish
8 under any authority or continue in effect as to any food in interstate commerce” followed
9 by a list of areas in which states may not enact laws in conflict with the federal laws.
10 These areas of food labeling which are expressly preempted fall under 21 U.S.C.A. §
11 343’s “Misbranded Foods” section. Included in this list are things such as offer for sale
12 under another name, imitation of another food, misleading container, etc.” Noticeably
13 absent from the express preemption section of 343-1, however, is “false or misleading
14 label.” Therefore, the Sherman Law, UCL, and FAL state law claims brought by
15 Plaintiffs, which are predicated on the basis that Defendant’s products’ labels are either
16 false or misleading, are not expressly preempted by the FDCA.

17 Next, Defendant contends that magnesium stearate, calcium stearate, and any other
18 stearate/stearic acid do not need to be labeled because 21 C.F.R. §101.100(a)(3)(ii) states
19 that incidental additives are exempt from labeling requirements if at insignificant levels.
20 (Def.’s Mot. Dismiss 10.) As Plaintiffs argue, however, whether these additives are
21 present in insignificant levels is a question of fact, not suited for dismissal at a preliminary
22 stage in the proceeding. The Court agrees and finds that plaintiffs’ claims are not expressly
23 preempted. Defendant’s motion to dismiss plaintiffs’ claims as expressly preempted will
24 be denied.

25 **2. Implied Preemption by the FDCA**

26 **i. Conflict with the FDCA Enforcement Scheme**

27 Defendant argues that even if Plaintiffs’ claims are not expressly preempted, they
28 are impliedly preempted because they conflict with the FDCA’s enforcement scheme.

(Def.’s Mot. Dismiss 12.) Such a conflict would show Congress’ intent for the federal government to occupy the field of food and beverage labeling. *Lockwood v. Conagra Foods, Inc*, 597 F.Supp.2d 1028 (N.D. Cal. 2009). This “intent may be inferred from a ‘scheme of federal regulation...so pervasive as to make reasonable the inference that Congress left no room for the States to supplement it,’ or where an Act of Congress ‘touch[es] a field in which the federal interest is so dominant that the federal system will be assumed to preclude enforcement of state laws on that subject.’” *English v. General Electric Co.*, 496 U.S. 72, 79 (1990) (citation omitted).

Defendant’s assertion that Plaintiffs’ claims conflict with the FDCA’s enforcement scheme is belied by the NLEA. The preemption provisions added to the FDCA by the NLEA include an express savings clause that disavows any implied preemption: “The NLEA shall not be construed to preempt any provision of State law, unless such provision is expressly preempted under 21 U.S.C. §343-1(a).” Pub.L. No. 101-535, §6(c)(1)(21 U.S.C. §343-1 note). “Congress has explicitly stated that it does not intend to occupy the field of food and beverage nutritional labeling; instead it permits states to regulate subject matters covered by the NLEA and its regulations provided that such state laws do not fall within the FDCA’s express preemption provisions.” *Lockwood* at 1032. Lastly, Defendant gives no explanation as to *how* Plaintiffs’ claims conflict with the FDCA’s enforcement scheme.

ii. Private Action Against the FDCA

Although citizens may petition the FDCA to take administrative action, private enforcement of the statute is barred. 21 U.S.C.A §337(a): “All such proceedings for the enforcement, or to restrain violations, of the Act shall be by and in the name of the United States.” *Id.*

Defendants allege that Plaintiffs’ claims are preempted because there is no private right of action to enforce FDA regulations. (Def.’s Mot. Dismiss 13.) Specifically, all claims for violations of the FDCA must be brought in the name of the United States. *Id.* Defendant argues that the FAC contains lengthy discussion of defendant’s alleged

1 violations of the FDA and various CFRs, thereby creating a private claim against the
2 FDA, which is prohibited.

3 Plaintiffs do not dispute that under the FDCA private litigants are expressly
4 prohibited from suing to enforce compliance with the federal regulations. However,
5 Plaintiffs contend that they are not attempting to enforce or to restrain violations of this
6 Act, but rather to enforce California's legal requirements, under the Sherman Law, UCL,
7 and FAL, which are identical to FDA regulations (Pl.s'. Opp'n at 13-14.)

8 Defendant's reliance in support of its argument on *Perez v. Nidek Co.*, 711 F.3d
9 1109 (9th Cir. 2013) is misplaced. In *Perez*, Plaintiffs claimed that Defendants failed to
10 disclose the lack of FDA approval for a specific type of laser required for LASIK surgery.
11 There, Plaintiffs brought suit with a claim for fraud by omission, something the court held
12 existed *solely* by virtue of the FDCA requirements. *Perez* at 1119. The court stated that
13 the existence of the federal enactments was a critical element in plaintiffs' case. *Id.* Those
14 facts are distinguishable from the instant case. Here, Plaintiffs' claims are not dependent
15 on the FDCA's existence. As Plaintiffs point out, the *Perez* court acknowledged that
16 plaintiff was "not barred from bringing *any* fraud claim related to the surgeries, [but] he
17 cannot bring a claim that rests solely on the non-disclosure to patients of facts tied to the
18 scope of . . . approval." *Id.* (emphasis included.)

19 In the present case, Plaintiffs are not suing because Defendant's conduct violates
20 the FDCA, rather, they are suing because Defendant's conduct allegedly violates
21 California's Sherman Law, UCL, and FAL, all of which "could have imposed the exact
22 same regulations even if the FDCA was never passed." *See Gustavson v. Wrigley Sales*
23 *Company*, 961 F.Supp.2d 1100, 1119 (N.D. Cal. 2013).

24 In light of the foregoing, Defendants motion on this basis will be denied.

25 **C. Sufficiency of Pleadings under Rules 8(a) and 9(b)**

26 Federal Rule of Civil Procedure 8(a) typically governs pleading requirements and
27 requires that a complaint contain "a short and plain statement of the claim showing that
28 the pleader is entitled to relief." FED. R. CIV. P. 8(a). However, a claim sounding in

1 fraud is subject to Rule 9(b)'s heightened pleading requirement that the plaintiff "state
 2 with particularity the circumstances constituting fraud." FED. R. CIV. P. 9(b). Rule 9(b)
 3 requires that allegations of fraud be "specific enough to give defendants notice of the
 4 particular misconduct . . . so that they can defend against the charge and not just deny that
 5 they have done anything wrong." *Vess v. Ciba-Geigy Corp. USA*, 317 F.3d 1097, 1106
 6 (9th Cir. 2003) (internal citations omitted). To satisfy Rule 9(b), the plaintiff must state
 7 the requisite "who, what, when, where, and how" of the misconduct he alleges. *See*
 8 *Kearns v. Ford Motor Co.*, 567 F.3d 1120, 1124 (internal citations omitted). The plaintiff
 9 must also "set forth an explanation as to why the statement or omission complained of
 10 was false and misleading when made," which can be done by "identifying either (1)
 11 inconsistent contemporaneous statements or (2) inconsistent contemporaneous
 12 information (such as an internal report) that was made by or was available to the
 13 defendants." *Yourish v. California Amplifier*, 191 F.3d 983, 993-94 (9th Cir. 1999).

14 Rule 9(b)'s heightened pleading standards apply to claims for violations of the
 15 UCL, FAL, and Sherman Act. *See Brazil v. Dole Food Co., Inc.*, 935 F.Supp.2d 947
 16 (N.D. Cal. 2013). Intentional misrepresentation and negligent misrepresentation are
 17 subject to the heightened pleading standards of Rule 9(b), which Plaintiffs do not dispute.
 18 *Rankine v. Roller Bearing Co. of America, Inc.*, 2013 WL 55802 (S.D. Cal. 2013).

19 **1. Standing to Bring Claims Regarding Unpurchased Products**

20 As addressed above, Plaintiffs lack standing to bring claims against Defendant's
 21 107 other products which they did not purchase. Therefore, this Court will not address the
 22 issue of sufficiency regarding those claims and instead will focus only on the sufficiency
 23 of claims brought against Defendant's One Daily Multivitamin and Men's One Daily.

24 **2. Claims Relating to Magnesium Stearate, Calcium Stearate, or any** 25 **other Stearate/Steric Acid**

26 Plaintiffs' FAC alleges that Defendant fraudulently omits the presence and use of
 27 "Magnesium Stearate, Calcium Stearate or any other Stearate/Stearic Acid" from its One
 28 Daily Multivitamin and Men's One Daily. (FAC ¶ 9, 42.) Defendant argues that this

1 allegation lacks the particularity required by Rule 9(b)'s heightened pleading standard.
2 (Def.'s Mot. Dismiss at 6.) Specifically, Defendant argues Plaintiff cannot show the
3 "who, what, when, where, and how" if Plaintiff cannot plead the specific additive they
4 allege exists within Defendant's products. Plaintiffs counter that they have sufficiently
5 alleged that Defendant has hidden and continues to hide the use of "Magnesium Stearate,
6 Calcium Stearate or any other Stearate/Stearic Acid" with the use of the term "vegetable
7 lubricant" instead of the common name as required by the FDA regulations. (Pl.s' Mot.
8 Dismiss at 7.) Plaintiffs then contend that Defendant chose to use this other name in order
9 to mislead consumers.

10 Plaintiffs have not fully met the heightened pleading standard. Plaintiffs' allegation
11 that Defendant's two products, One Daily Multivitamin and Men's One Daily, contain
12 "Magnesium Stearate, Calcium Stearate or any other Stearate/Stearic Acid" fails to meet
13 9(b)'s heightened standard because it does not "give defendants notice of the particular
14 misconduct which is alleged to constitute the fraud charge[s]." *Brazil v. Dole Food Co.,*
15 *Inc.*, 935 F.Supp.2d 947 (N.D. Cal. 2013). Plaintiffs will need to specify which additives
16 they are claiming are omitted from Defendant's products in order to meet 9(b)'s standard.
17 It is not enough to simply state "or any other Stearate/Stearic Acid," as this does not give
18 requisite notice to Defendant in order for it to defend claims against it.

19 In *Brazil v. Dole Food Co., Inc.*, 935 F.Supp.2d 947 (N.D. Cal. 2013) the court
20 determined that the plaintiff's pleadings failed to meet the heightened pleading
21 requirement because 1) it failed to specify which products were at issue in the case; 2) no
22 precise nature of any alleged violations was stated; and 3) it did not specify which
23 content on the labels plaintiff had relied upon and found misleading. *See Brazil v. Dole*
24 *Food Co., Inc.*, 935 F.Supp.2d at 964. Plaintiffs in this case, in contrast to *Brazil*, have
25 sufficiently plead the who, what, when, where, and how. They have stated the time period
26 in which the alleged behavior ensued, described the alleged mislabeling at great length,
27 with photos of the products' labels included, and argued why Defendant allegedly
28 mislabels its products. (FAC ¶ 44, 46.)

1 Plaintiffs will be given leave to amend this claim with further facts concerning
2 “any other stearate/stearic acid.”

3 **3. Plaintiffs’ Have Plead their Claims Regarding Defendant’s**

4 **Misrepresentation of Whole Foods with Sufficient Particularity**

5 Defendant argues that all of Plaintiffs’ claims should be dismissed regarding the
6 whole food content of Defendant’s products, One Daily Multivitamin and Men’s One
7 Daily. In support of this argument, Defendant contends that Plaintiffs make vague claims
8 such as the products containing “much less [whole foods] than customers reasonably
9 expect” and that it is “*likely* that the products they bought did not provide the stated
10 amounts.” (Def.’s Mot. Dismiss at 7.)

11 Plaintiffs have sufficiently pleaded facts that the products do not contain vitamins
12 and minerals strictly from whole food sources. Even if synthetically-produced minerals
13 and vitamins were present at extremely low or negligible levels, Plaintiffs argument is
14 sufficient to meet 9(b) because they contend that they were misled by a representation
15 that *only* whole food sources were utilized. Any amount of synthetically-produced
16 minerals or vitamins would be a misrepresentation according to this line of reasoning.

17 Further, Plaintiff alleges numerous times how they were misinformed and that
18 “synthetic or processed nutrients were added to [Defendant’s] products. (FAC ¶ 23.)
19 Plaintiffs’ basis for this argument is that “it is impossible to provide the amount of
20 vitamins and minerals from the amount of whole foods that Defendant purports to use to
21 obtain the levels of vitamins and minerals as labeled on its packages.” (FAC ¶ 41.)

22 Finally, Defendant argues that Plaintiff should state facts about what the product
23 actually contains. (Def.’s Mot. Dismiss at 7.) Plaintiffs do just this. Plaintiffs state there
24 are synthetic, as well as whole food, sources of nutrients and minerals in the products.
25 (FAC ¶ 94, 101, 109.)

26 In light of the foregoing, Defendants motion on the basis of failure to plead fraud
27 with particularity will be granted in part and denied in part. Plaintiffs may amend the
28 complaint to correct the deficiency noted.

